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Radhakrishnan, Anand N. P.; Pradas, Marc; Sorensen, Eva; Kalliadasis, Serafim and Gavriilidis, Asterios (2019). Hydrodynamic Characterization of Phase Separation in Devices with Microfabricated Capillaries. *Langmuir*, 35(25) pp. 8183–8516.

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Version: Version of Record

Link(s) to article on publisher's website:

<http://dx.doi.org/doi:10.1021/acs.langmuir.8b04202>

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Hydrodynamic Characterization of Phase Separation in Devices with Microfabricated Capillaries

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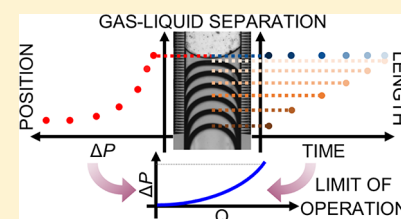
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S Supporting Information

ABSTRACT: Capillary microseparators have been gaining interest in downstream unit operations, especially for pharmaceutical, space, and nuclear applications, offering efficient separation of two-phase flows. In this work, a detailed analysis of the dynamics of gas–liquid separation at the single meniscus level helped to formulate a model to map the operability region of microseparation devices. A water–nitrogen segmented flow was separated in a microfabricated silicon–glass device, with a main channel (width, $W = 600\ \mu\text{m}$; height, $H = 120\ \mu\text{m}$) leading into an array of 276 capillaries ($100\ \mu\text{m}$ long; width = $5\ \mu\text{m}$ facing the main channel and $25\ \mu\text{m}$ facing the liquid outlet), on both sides of the channel. At optimal pressure differences, the wetting phase (water) flowed through the capillaries into the liquid outlet, whereas the nonwetting phase (nitrogen) flowed past the capillaries into the gas outlet. A high-speed imaging methodology aided by computational analysis was used to quantify the length of the liquid slugs and their positions in the separation zone. It was observed that during stable separation, the position of the leading edge of the liquid slugs (advancing meniscus), which became stationary in the separation zone, was dependent only on the outlet pressure difference. The trailing edge of the liquid slugs (receding meniscus) approached the advancing meniscus at a constant speed, thus leading to a linear decrease of the liquid slug length. Close to the liquid-to-gas breakthrough point, that is, when water exited through the gas outlet, the advancing meniscus was no longer stationary, and the slug lengths decreased exponentially. The rates of decrease of the liquid slug length during separation were accurately estimated by the model, and the calculated liquid-to-gas breakthrough pressures agreed with experimental measurements.



INTRODUCTION

Capillary-based separators are currently seeing a significant growth in their implementation for downstream separation of two-phase flows. Capillarity-driven separation can be exploited for their high efficiency and accurate spatiotemporal control afforded by advancements in microfluidics.^{1,2} Much of its application has been in microchemical reaction systems,³ where, for example, a gaseous component in a reaction requires a gas–liquid separation step to isolate the two phases. Such devices have also been applied for liquid–liquid extraction,^{4–10} gas–liquid separation in fuel cells,¹¹ and safe handling of toxic or carcinogenic molecules.¹² Capillarity-driven separators utilize either membranes or microfabricated capillaries to separate the phases. For membrane-based separators,^{13–16} it has been observed that factors such as membrane area, thickness, wettability, pore size, and tortuosity affect the separation process. Microfabricated separators^{17,18} achieve separation similarly by taking advantage of the differences in the wettability between the two phases and the surface tension of the liquid, with the added advantage that the pores are microengineered, hence having a uniform size, and are not tortuous. Although limited in their throughput, the fabrication

of uniform capillary structures facilitates selective separation with controllable system behavior. For a typical hydrophilic substrate (e.g., silicon or glass), the wetted phase (e.g., water) flows through these capillaries, whereas the nonwetting dispersed phase (gas) is prevented from entering the capillaries due to the capillary pressure.^{19,20} Liquid–liquid separation can also be achieved through a similar methodology²¹ by taking advantage of the differences in the wetting properties of the two liquids.

The underlying phenomena of phase separation in capillary microseparators, at a scale where interfacial tension and capillary effects are dominant, are far from being fully understood. In terms of operational limits of capillary microseparators, the pressure at which the gas phase enters the capillaries (gas-to-liquid breakthrough), $\Delta P_{G>L}$, can be estimated from the Young–Laplace equation (appropriately modified for rectangular channels²²)

Received: December 18, 2018

Revised: May 15, 2019

$$\Delta P_{G>L} = P_G - P_L = 2\gamma \cos \theta_e \left(\frac{1}{D} + \frac{1}{H} \right) \quad (1)$$

where γ is the surface tension, D and H are the capillary width and height (reflecting the principle radii of curvature of the meniscus), respectively, θ_e is the equilibrium contact angle of the wetting liquid on a substrate, and P_G and P_L are the gas outlet and liquid outlet pressures, respectively. On the other hand, the liquid-to-gas breakthrough pressure, $\Delta P_{L>G}$, (i.e., the point at which complete liquid separation ceases and liquid exits through the gas outlet) is typically estimated from the pressure drop across the capillaries in the microseparator using the Hagen–Poiseuille equation. Several groups have reported deviations from these estimates for both gas–liquid and liquid–liquid separations. Various parameters were then included in the breakthrough models^{15,19,21,23,24} to account for (i) flow of liquid through N capillaries in parallel, (ii) geometric correction factors (e.g., as described by Mortensen et al.²⁵), (iii) variations in apparent contact angles of the menisci, and (iv) the liquid slug lengths and number of active capillaries during separation. This modified equation is commonly given as

$$\Delta P_{L>G} = P_G - P_L = \frac{\alpha \eta L_c}{N A_c^2} Q_T \quad (2)$$

where Q_T is the total inlet flow rate assumed to be equal to NQ_c . Here, Q_c is the flow through a single capillary, η is the viscosity, L_c is the length of the capillaries, A_c is the cross-sectional area of the capillaries, and α is a correction factor that is yet to be fully characterized.

To estimate the local flow rate through each capillary (Q_c), knowledge of the pressure drop across them is required. This pressure drop is often estimated from the pressure difference between the local pressure at the entrance of a capillary and the liquid-side outlet pressure, assuming that all N capillaries are used for separation (i.e., leading to $Q_T = NQ_c$). There is, however, enough experimental evidence suggesting that not all capillaries are used for separation.²⁶ Furthermore, we are yet to understand and characterize the effect of capillary geometry, and hence, a geometric correction factor is still used in the Hagen–Poiseuille equation (e.g., α in eq 2). In the literature, only empirical equations exist to estimate the correction factor for rectangular or nonrectangular cross sections^{25,27} that are constant along the length of the capillary. However, in devices with nonparallel capillary walls, the cross-sectional geometry varies along the length of the capillary, thus affecting the local pressure drop,^{17,19} and such empirical correction factors can no longer be applied. Experimental investigation on such a capillary design could yield crucial information not only regarding the flow rate of the liquid through the Hagen–Poiseuille equation but also regarding the wetting properties in the capillaries and thus the gas-to-liquid breakthrough pressures. Experimental measurement of the contact angles in the capillaries is challenging due to limitations in optical imaging (i.e., imaging needs to be performed inside high aspect ratio capillaries with narrow widths and high depths, where the reflection of light is poor). Preliminary attempts with fluorescence microscopy were performed by our group to measure the velocity inside the capillaries. However, the presence of fluorophores in the liquid under investigation led to changes in surface tension, thus affecting the separation process. As regards the length of the liquid slugs in the segmented flow, which influences the separation process,^{19,21}

investigations have been performed to either experimentally measure the slug lengths upstream of the capillary zone or estimate them from scaling laws (e.g., the model proposed by Garstecki et al.²⁸). However, the estimation from the scaling law by Garstecki et al. does not account for the dynamic variations in the liquid slug length during separation. This is because the formation of liquid slugs upstream of the capillaries is affected by the pressure variations due to the balancing of forces between the net inflow of liquid into the separation zone and the outflow of liquid through the capillaries.²³ Additionally, in gas–liquid separation, the variation in the gas bubble length due to the pressure drop across the two-phase flow in the inlet segment also leads to variations in the liquid slug lengths before entering the separation zone. All model estimations in the literature eventually result in a linear behavior of the liquid-to-gas breakthrough pressure ($\Delta P_{L>G}$) with respect to liquid flow rate, where the complex hydrodynamics surrounding the two-phase flow remains unresolved.

In previous work, we experimentally investigated the effect of flow rates of a gas–liquid-segmented flow in a capillary microseparation device¹⁹ and also undertook a theoretical analysis of the breakthrough limits. In the present work, we advance the understanding of gas–liquid separation by developing a hydrodynamic model based on a liquid slug during separation in an array of microcapillaries. The development of the model is aided by *in situ* analysis of the liquid slug properties resolved by high-speed imaging, combined with a computational image analysis algorithm.

■ EXPERIMENTAL SETUP

Microfluidic Device Fabrication. Two gas–liquid separator designs, A-1 and A-2, with trapezoidal-shaped capillaries with a rectangular cross section (see Table S1 and Figure S1 in the Supporting Information for the design parameters) were fabricated on a 4 in. silicon (Si) wafer (PI-KEM Limited, U.K.) via photolithography and deep reactive ion etching (DRIE) as described by Roydhouse et al.¹⁹ The silicon wafer was spin-coated with a positive photoresist (SPR-220-7, Rohm and Haas, USA) and soft-baked at 90 °C for 2 min, followed by 115 °C for 3 min. The separator pattern was transferred onto the photoresist using a UV mask aligner (Q4000-6, Quintel, USA). Pattern development was performed by immersing the wafer in a photoresist developer (MF-26A, Shipley Microposit, Dow Chemical Company, USA) for 2.5 min followed by thorough rinsing with deionized water. The wafer was placed at 115 °C for 3 min as a post-exposure baking step. The device was then etched using a DRIE instrument (ICP DRIE, SPTS Technologies, U.K.) with a standard Bosch process. The depth of the etched devices was measured using a surface profiler (DektakXT, Bruker, U.K.). The wafer was diced into the desired dimensions using a precision dicing saw (DAD 3230, DISCO Corporation, Japan). A glass slide (Corning 7740, Newcastle Optical Engineering, U.K.), with predrilled holes for fluidic inlet and outlets, was anodically bonded to the Si wafer (400 °C, 500 VDC, 5 min). Before bonding, the substrates were cleaned with acetone and deionized water (Millipore grade), followed by a final O₂ plasma clean (Plasma Asher, Diener electronic, Germany) for 10 min.

Gas–Liquid Separation Procedure. The separation performance was studied experimentally on a water–nitrogen system using the setup shown in Figure 1. Deionized water (Millipore grade, surface tension = 0.07 N/m, density = 1000 kg/m³, viscosity = 8.9 × 10^{−4} Pa·s) was procured in-house and injected into a T-junction (internal diameter = 1 mm, Upchurch Scientific, USA) using a displacement syringe pump (Legato 270P, KD Scientific, USA) at varying flow rates (Q_L). Zero grade nitrogen (BOC, U.K.) was introduced via a mass flow controller (Bronkhorst, Netherlands),

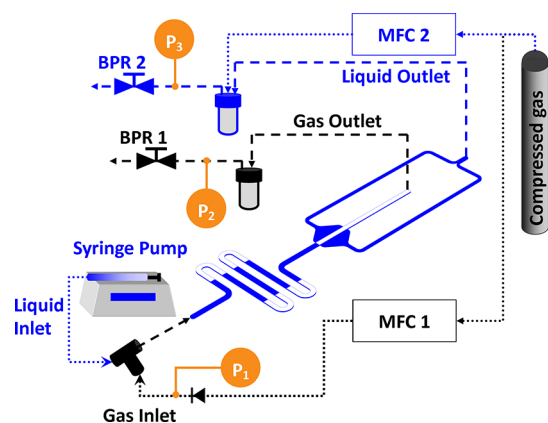


Figure 1. Schematic representation of the experimental setup. P_1 – P_3 represent points of pressure monitoring. BPR: back pressure regulator; MFC: mass flow controller (image not to scale).

MFC 1 (in Figure 1), into the second port of the T-junction at a constant flow rate Q_G of 350 $\mu\text{L}_n/\text{min}$ to achieve a gas–liquid-segmented flow, which was introduced into the device. Pressures were recorded using 0–15 psi pressure sensors (40PC015G Honeywell, U.K.), shown as P_1 – P_3 in Figure 1, connected to a data acquisition board (USB-6000, National Instruments, U.K.) monitored via a script written on LabView 2015 (National Instruments, U.K.). The pressures at the gas and liquid outlets were controlled using adjustable back pressure regulators (KCB Series, Swagelok, U.K.) of ranges 0–10 psi (BPR 1 in Figure 1) and 0–25 psi (BPR 2 in Figure 1), respectively. The gas outlet was connected to a low-volume liquid collection vial (2 mL glass vial, Agilent Technologies, U.K.) before BPR 1 to trap any liquid exiting the gas outlet, especially during liquid-to-gas breakthrough case studies. Similarly, the liquid outlet was connected to a vessel (100 mL borosilicate glass, VWR International, U.K.) to collect the liquid exiting the device before BPR 2. The liquid collection vessel was pressurized by a separate nitrogen line at 20 mL_n/min controlled by MFC 2.

The operating procedure followed to measure the breakthrough pressures is described in Figure S2 of the Supporting Information. The P_2 pressure sensor recorded the gas outlet pressure (P_G), whereas the P_3 sensor recorded the liquid outlet pressure (P_L). To establish successful separation, P_G was kept higher than P_L , that is, the pressure difference ΔP_{GL} was always positive. Considering a pure gas flow, the pressure drop from the last capillary at the gas outlet in the device to the pressure sensor was estimated to be around 4.2 Pa using the Hagen–Poiseuille equation, whereas the maximum pressure drop of water from the exit of the capillaries to the pressure monitoring point was around 302 Pa.

High-Speed Imaging and Computational Image Analysis.

The separation and breakthrough experiments were monitored using a high-speed camera (Mini AX100, Photron, USA) mounted on a microscope (Axioscope A1, Carl Zeiss, U.K.) with a vibration

isolation platform (Newport Spectra-Physics, U.K.). Images were captured at 4000 frames/s for liquid flow rates (Q_L) above 30 $\mu\text{L}/\text{min}$ and at 1800 frames/s for Q_L below 30 $\mu\text{L}/\text{min}$ at 10 μs exposure. The images (saved as TIFF) were processed offline. An automated image analysis script was developed in the Python programming language^{29–33} to measure the slug length and number of capillaries used by the liquid slug in the separation zone. Upon execution of the script, a sequence of images was opened, and they were then converted to a pixel array. The only manual intervention required from the user was to locate the walls of the main channel and the position of the first capillary on the images. The corresponding pixel coordinates were then used to measure the length scales in the images. To remove noise and small water droplets left on the top wall or the bottom of the separation zone, an image reconstruction step was adopted based on a morphological erosion filter (see Figure S3 of the Supporting Information). The reconstructed pixel arrays were then binarized to a Boolean array. Different contours in the binary images were estimated by an edge detection algorithm followed by a gradient filter. Only the contours touching both the edges of the walls were chosen as the menisci of the gas–liquid slugs, and the distance as well as the number of capillaries between them were measured (see Supplementary Movie 1).

THEORETICAL FRAMEWORK

A hydrodynamic theory for the calculation of the liquid-to-gas breakthrough pressure, $\Delta P_{L>G}$, calculation is developed next. We assume that the system is isothermal at room temperature and that the viscosities and densities of the liquid and gas phases remain constant during the experiment. We consider a liquid slug of initial length (L_0) moving along the main channel of the microseparator at the superficial speed $U = Q_T/A$, where A is the cross-sectional area of the main channel and $Q_T = Q_G + Q_L$ is the total flow rate that is the sum of the liquid and the gas flow rates. When the slug enters the separation zone (i.e., the array of capillaries), we consider the entrance of the separation zone as the reference point ($X = 0$) and denote the position of the advancing meniscus with respect to this origin as $X_a(t)$ and the position of the receding meniscus as $X_r(t)$ (see Figure 2). We consider that the advancing meniscus reaches the origin, that is, the first capillary, at time $T = 0$, such that $X_a(0) = 0$. In the following, we consider two different stages, namely, stage I in which the separator achieves stable separation and stage II for which the separator approaches critical breakthrough conditions. As shown below, the crucial difference between these two stages is that during the stage I, the advancing interface is stationary, whereas the receding interface is moving. On the other hand, during the stage II, both interfaces are moving toward the gas outlet.

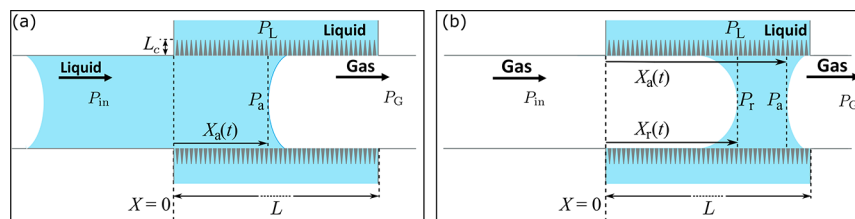


Figure 2. Schematic representation of gas–liquid separation in the microseparator. (a) A liquid slug enters the separation zone of length L , and we denote the position of the advancing interface with respect to the first capillary as $X_a(t)$. The liquid flows through the capillaries of length L_c . (b) When the receding interface enters the separation zone, its position is denoted as $X_r(t)$. P_a is the liquid-side pressure at the advancing interface, and P_r is the liquid-side pressure at the receding interface. P_{in} is the total inlet pressure, whereas P_G and P_L are the gas outlet and liquid outlet pressures, respectively, controlled by the user.

Stage I: Stable Separation. As the slug enters the separation zone, we denote the mass of the part of the slug in contact with the microcapillaries as M , which is given by

$$M(T) = \rho A X_a(T) \quad (3)$$

where ρ is the density of the liquid. This mass changes over time due to the inflow of liquid that enters the separation zone, which is given by $Q_L \rho$, and the outflow of liquid through the capillaries (M_{outflow}), given by

$$\frac{dM_{\text{outflow}}(T)}{dT} = Q_c \rho N \frac{X_a(T)}{L} \quad (4)$$

where Q_c is the local flow rate through each capillary, N is the total number of capillaries, and L is the length of the separation zone (see Figure 2). During stable separation, as not all capillaries are available for separation, M_{outflow} is the flow through the total number of capillaries (N) scaled by the position of the advancing meniscus in the separation zone (X_a) with regards to the total length of the separation zone, L . From a mass balance of the liquid slug undergoing separation under these inflow and outflow conditions as given in eq 4, we get

$$\frac{dM(T)}{dT} = Q_L \rho - Q_c \rho N \frac{X_a(T)}{L} \quad (5)$$

Substituting eq 3 in eq 5 and rearranging for X_a , we find

$$\frac{dX_a}{dT} = \frac{Q_L}{A} - Q_c \frac{N}{AL} X_a \quad (6)$$

We now estimate the local flow rate, Q_c , at each capillary by making use of a local Hagen–Poiseuille law

$$Q_c = \frac{A_c^2}{\alpha \eta L_c} \Delta P_c \quad (7)$$

where η is the viscosity, L_c is the length of the capillaries, and α is a geometrical correction for a single capillary (refer to the works of Mortensen et al.²⁵ or Bahrami et al.²⁷). The pressure difference ΔP_c is given by the pressure drop along each capillary: $\Delta P_c = P_a - P_L$, where P_a is the pressure at the advancing interface and can be estimated as $P_a = P_G - P_c$, so that

$$\Delta P_c = P_G - P_L - P_c = \Delta P_{GL} - P_c \quad (8)$$

Here, P_c is the capillary pressure of the advancing interface in the main channel, calculated from the Young–Laplace equation similar to the form shown in eq 1. $\Delta P_{GL} = P_G - P_L$ is the controlled pressure drop of the microseparator.

We introduce nondimensional variables by taking the length of the separation zone, L as the length scale, $T_0 = L/U_0$ as the time scale, and $P_0 = \eta U_0/L_c$ as the pressure scale, where $U_0 = Q_G/A$ is the typical speed given by the constant gas flow rate Q_G . With these considerations, the differential eq 6 becomes

$$\frac{dx_a}{dt} = q_L - \beta \Delta p x_a \quad (9)$$

where we have defined $x_a = X_a/L$, the dimensionless pressure drop $\Delta p = (\Delta P_{GL} - P_c)/P_0$, the dimensionless time $t = T/T_0$, the dimensionless liquid flow rate $q_L = Q_L/Q_G$, and the parameter

$$\beta = NA_c^2/\alpha AL_c^2 \quad (10)$$

Solving eq 9 with the condition $x_a(0) = 0$, we obtain

$$x_a(t) = \frac{q_L}{\beta \Delta p} (1 - e^{-\beta \Delta p t}) \quad (11)$$

On the other hand, the position of the receding interface, X_r , can be described as $X_r = UT - L_0$, based on the gas bubble moving at a constant speed, $U = Q_T/A$. In dimensionless units, where $x_r = X_r/L$, we have

$$x_r(t) = q_T t - l_0 \quad (12)$$

where $l_0 = L_0/L$, $q_T = Q_T/Q_G$, and L_0 is the initial length of the liquid slugs. We note that we do not consider any slip effects due to thin liquid films on the wall. Because we are only interested in the dynamics of the liquid slug inside the separation zone, any possible effect of slip on the gas bubble velocity can be neglected as any liquid along the walls is imbibed through the capillaries. As shown below, our results indicate that this is a reasonable assumption. When the receding interface enters the separation zone, that is, $x_r(t) = 0$ in eq 12, we get

$$t = l_0/q_T \quad (13)$$

At this time t , the advancing interface is at location x_p , obtained by substituting eq 13 in eq 11:

$$x_p = \frac{q_L}{\beta \Delta p} (1 - e^{-\beta \Delta p l_0/q_T}) \quad (14)$$

In the stage I considered here for which separation is stable, we assume that the advancing meniscus is always pinned, that is, the advancing meniscus is stationary with x_p constant, after the receding meniscus has entered the separation zone, as reported experimentally in other studies.^{17,19} Note that x_p is always <1 and at $t = l_0/q_T$, the receding meniscus enters the separation zone, that is, $x_r(l_0/q_T) = 0$, and hence, the dimensionless length of the slug is at this point $l_s(l_0/q_T) = x_a - x_r = x_p$. Because the receding interface moves at the constant speed of $q_T t$, the length of the slug decreases linearly as

$$l_s(t) = x_p - q_T t \quad (15)$$

where we have considered that $t = 0$ is the time when the receding interface enters the separation zone.

Stage II: Critical Conditions. For a given flow rate, as the pressure difference ΔP_{GL} is decreased, the advancing meniscus may not reach a pinned position before the receding meniscus enters the separation zone. Under these conditions, the variation of the liquid slug length is not linear, and eq 15 needs to be modified to account for the dynamics of the advancing interface. In particular, we note that the outflow of liquid through the capillaries as in eq 4 is now given by

$$\frac{dM_{\text{outflow}}(T)}{dT} = Q_c \rho N \frac{X_a(T) - X_r(T)}{L} \quad (16)$$

where $X_r(T)$ is the location of the receding interface. Following the same mass balance as in eq 5 to obtain eq 9, we now arrive at

$$\frac{dx_a}{dt} = q_L - \beta \Delta p (x_a - x_r) \quad (17)$$

$$\frac{dx_r}{dt} = q_T \quad (18)$$

Equations 17 and 18 can be combined to get

$$\frac{dl_s}{dt} + \beta \Delta p l_s = -1 \quad (19)$$

where, $l_s(t) = x_a - x_r$, $q_L - q_T = -1$, and $\Delta p = (\Delta P_{GL} - P_c)/P_0$, as defined above. We note that the location x_p given by eq 14 corresponds to the location of the advancing interface when the receding interface enters the separation zone. Therefore, we solve the differential eq 19 with the condition $l_s(0) = x_p$ to obtain

$$l_s(t) = \left(x_p + \frac{1}{\beta \Delta p} \right) e^{-\beta \Delta p t} - \frac{1}{\beta \Delta p} \quad (20)$$

For a given flow rate, the liquid-to-gas breakthrough occurs when $l_s = 0$ and $x_r = 1$, namely, when the slug length is zero and at the same time, the receding meniscus is at the exit of the separation zone. Because $x_r(t) = q_T t$, this condition is given by

$$\Delta p_{L>G} = \frac{1}{\beta} q_T \ln[1 + q_L(1 - e^{-\beta \Delta p_{L>G} l_0/q_T})] \quad (21)$$

where we have used eq 14. If the initial length of the slugs, l_0 , is sufficiently large, we can approximate $e^{-\beta \Delta p_{L>G} l_0/q_T} \approx 0$, and so we have

$$\Delta p_{L>G} \approx \frac{1}{\beta} q_T \ln q_T \quad (22)$$

We note that as expected, if the liquid flow rate is zero, $q_L = 0$, we then have $q_T = 1$ and $\Delta p_{L>G} = 0$. The developed model from eqs 21 and 22 captures the dynamics of a liquid slug during separation and yields a nonlinear behavior of the liquid-to-gas breakthrough pressures, as opposed to a linear estimation from eq 2, which has been proposed so far in the literature.^{15,19,21,23} This breakthrough pressure can be dimensionalized with P_0 and can be expressed as

$$\Delta p_{L>G} \approx \frac{P_0}{\beta} \frac{Q_T}{Q_G} \ln \frac{Q_T}{Q_G} + P_c \quad (23)$$

RESULTS AND DISCUSSION

Initial Observations of Stable Separation. The separation of the water–nitrogen-segmented flow was tested in the two microseparation devices, A-1 and A-2 (see Table S1 and Figure S1 in the Supporting Information for the design parameters), at varying liquid flow rates (Q_L) and a constant gas flow rate Q_G of 350 $\mu\text{L}_n/\text{min}$. The two devices had different capillary geometry and number of capillaries. Capillaries with a tapered geometry, as first reported by Günther et al.,¹⁷ have a higher throughput in comparison with rectangular geometries. Although the tapered capillaries provide comparable $\Delta P_{G>L}$ values as that of rectangular capillaries, the $\Delta P_{L>G}$ values are lowered. For a given pressure drop across each capillary (ΔP_c), Q_c is higher in tapered capillaries due to a larger cross-sectional area (A_c) and thus the operating zone of such microseparation devices is improved.¹⁹ The pressure difference (ΔP_{GL}), which is the difference between the gas outlet pressure (P_G at P_2 in Figure 1) and the liquid outlet pressure (P_L at P_3 in Figure 1), was controlled with the help of BPRs and monitored using pressure sensors connected to LabView via a data acquisition board. The pressure drop between the separation zone and the pressure monitoring points at the outlets was considered to be negligible, as shown in the Gas–Liquid Separation Procedure

section. For all experiments performed, ΔP_{GL} was first set at a suitable value at which stable separation occurred using only gas flow (nitrogen) followed by the injection of water (to initiate segmented flow). The capillaries were, therefore, prewetted by the water in the slug flow to avoid the collection of the dispersed phase (nitrogen) inside the capillaries. This is because capillary prewetting, which is often overlooked, is an important factor that affects the capillary pressures and thus the operability of the device. Once the device was appropriately prewetted with water (the wetting phase), the capillaries were observed to be filled with water, and the flow through the capillaries during separation was pressure driven as described by the Hagen–Poiseuille law (eq 7). Gas–liquid separation in the A-1 device was observed using high-speed imaging as shown in Figure 3. The gas bubble approaching the

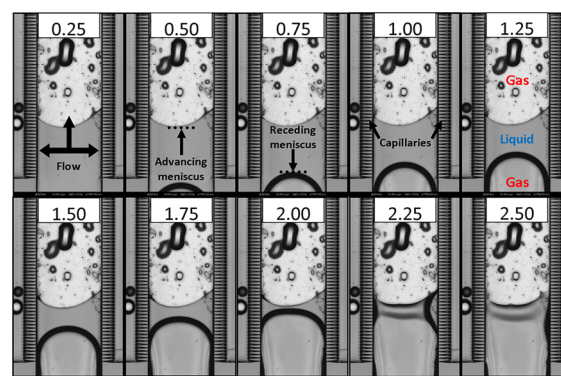


Figure 3. Sequence of high-speed images of a liquid slug in the separation zone during stable separation in the A-1 device, with gas flow rate $Q_G = 350 \mu\text{L}_n/\text{min}$ and liquid flow rate $Q_L = 90 \mu\text{L}/\text{min}$ at outlet pressure difference $\Delta P_{GL} = 7 \text{ kPa}$. The advancing meniscus is observed to be pinned as all the liquid exits through the capillaries. The inset labels represent the observation time in milliseconds. The flow is from the bottom to top. Images were recorded at 4000 frames/s.

advancing meniscus was covered by a liquid film, whereas the gas phase toward the gas outlet was devoid of it (see Figures 3 and 4). This was because the trailing film of the preceding gas bubble had sufficient time to coalesce into water droplets as can be seen in the main channel (also see Supplementary Movie 2 for the entire sequence of water separating at $Q_L = 90 \mu\text{L}/\text{min}$ and the immediate coalescence of the liquid film into water droplets in the channel). These droplets were consequently filtered out from the images so as to accurately identify only the liquid slug menisci. At stable separation conditions, the advancing meniscus was observed to be pinned inside the separation zone.

In the first instance, high-speed images were taken at the entrance of the A-1 device (upstream of the separation zone) during stable separation, and the initial length of the slugs, l_0 , and number of capillaries used for separation were measured via ImageJ.³⁴ It was observed that the number of capillaries that were active during separation increased with decreasing ΔP_{GL} (Figure S4c of the Supporting Information) because the advancing meniscus moved further inside the separation zone as ΔP_{GL} approached the liquid-to-gas breakthrough pressures ($\Delta P_{L>G}$). On the other hand, the number of active capillaries increased with increasing Q_L (Figure 4 and Figure S4d of the Supporting Information).

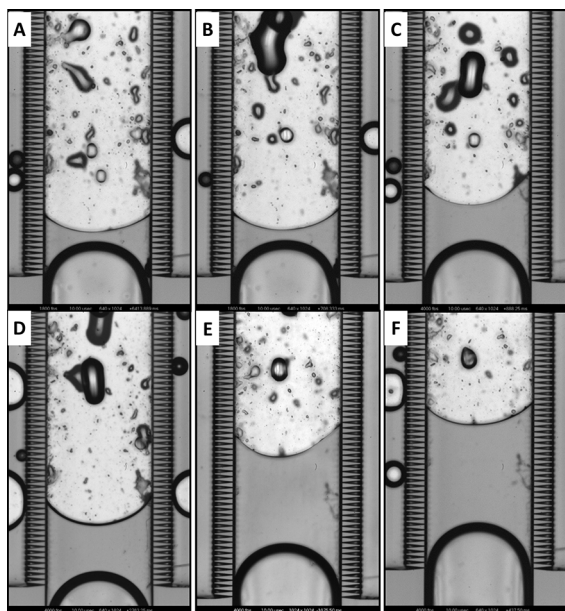


Figure 4. Snapshots of high-speed images showing the position of the advancing meniscus inside the separation zone, as a function of liquid flow rate, during stable separation in the A-1 device. Gas flow rate $Q_G = 350 \mu\text{L}_n/\text{min}$ with varying liquid flow rates $Q_L =$ (A) $9 \mu\text{L}/\text{min}$, (B) $20 \mu\text{L}/\text{min}$, (C) $90 \mu\text{L}/\text{min}$, (D) $120 \mu\text{L}/\text{min}$, (E) $150 \mu\text{L}/\text{min}$, and (F) $200 \mu\text{L}/\text{min}$ at outlet pressure difference $\Delta P_{GL} = 7 \text{ kPa}$. The flow is from the bottom to top. Images were recorded at 4000 frames/s.

A significant fluctuation in the pinning positions, however, was observed, especially as the gas-to-liquid and liquid-to-gas breakthrough pressures converged at high Q_L , thus reducing the operability window. Accurate control of the pressure difference, which affects the pinning position of the advancing menisci of the liquid slugs, became progressively harder even with the use of low-range, high-sensitivity BPRs. At high flow rates, where the net inflow of liquid into the separation zone was high, the pressure fluctuations were caused when the advancing meniscus instantaneously became stationary. Because of this highly dynamic nature of separation, small gas bubbles were observed to enter the liquid collection ducts through the capillaries due to local instabilities in pressure as

seen on the left side of the images in Figures 3 and 4. These gas bubbles were not observed to affect the separation, and the occurrence of such events was sporadic and at random locations in the capillary array. The liquid in the liquid collection ducts was considered to have a uniform pressure drop until the outlets of the device, as it was observed to be devoid of any gas bubbles.

The initial slug lengths, L_0 , which influence the liquid-to-gas breakthrough pressures,^{19,21} seemed fairly constant at around 70 mm at varying ΔP_{GL} , where stable separation was observed (Figure S4a of the Supporting Information). The length of slugs was measured during the process of separation with a significant pressure drop across the main channel along with the aforementioned pressure fluctuations. The formation of slug flow depends on the pressure difference between the continuous and dispersed phases, as well as on the interfacial tension, and is affected by the pressure changes at the separation zone.²³ During the slug formation at the T-junction, the dispersed bubble enters the main stream and elongates. During this phase, before the bubble pinches-off from the dispersed phase inlet, the continuous phase pressure increases and pushes more fluid through the film around the dispersed bubble, whereas the dispersed phase pressure remains constant. During stable separation, the pinning of the advancing meniscus of the liquid slug in the separation zone creates an instantaneous increase in the slug flow pressure upstream. Additionally, the two-phase flow separates into two single-phase flows in the separation zone, with the continuous phase bifurcating into two liquid outlet streams, leading to a varying pressure field. Furthermore, the difference between P_{in} and P_G , which is directly proportional to Q_L , ranged between 3–17 kPa (as seen in Figure S2 of the Supporting Information). At these pressure drops, the gas bubble is expected to expand gradually, and consequently, the local velocity of the liquid slug increases. This in turn leads to a decrease in the liquid slug length along the length of the main channel before reaching the capillary array. Under such conditions, for an accurate estimation of liquid slug lengths (L_0) in systems with significant pressure drops, the readers can refer to the work reported by Molla et al.³⁵ alongside the scaling law proposed by Garstecki et al.²⁸

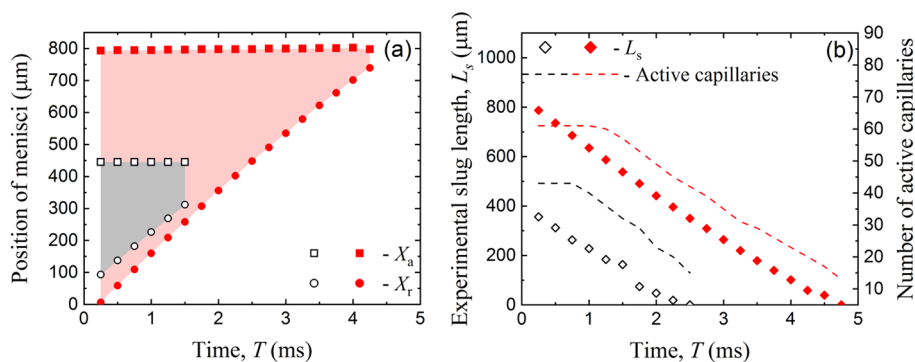


Figure 5. Tracking of liquid slugs during stable separation in the A-1 device. (a) Positions of the advancing (square markers, X_a) and receding (circle markers, X_r) menisci from the first capillary. The graph shows the linear motion of the receding meniscus toward the pinned advancing meniscus as liquid separates through the capillaries. Liquid flow rate $Q_L = 90 \mu\text{L}/\text{min}$ (black, open markers) and $200 \mu\text{L}/\text{min}$ (red, closed markers) at gas flow rate $Q_G = 350 \mu\text{L}_n/\text{min}$ and outlet pressure difference $\Delta P_{GL} = 7.03 \text{ kPa}$. (b) The diamond markers show the length of the liquid slugs ($L_s = X_a - X_r$) within the separation zone measured between the meniscus caps. Dashed lines denote the number of active capillaries as the receding meniscus enters the separation zone. $Q_L = 90 \mu\text{L}/\text{min}$ (black, open markers) and $200 \mu\text{L}/\text{min}$ (red, closed markers) at $Q_G = 350 \mu\text{L}_n/\text{min}$ and $\Delta P_{GL} = 7.03 \text{ kPa}$.

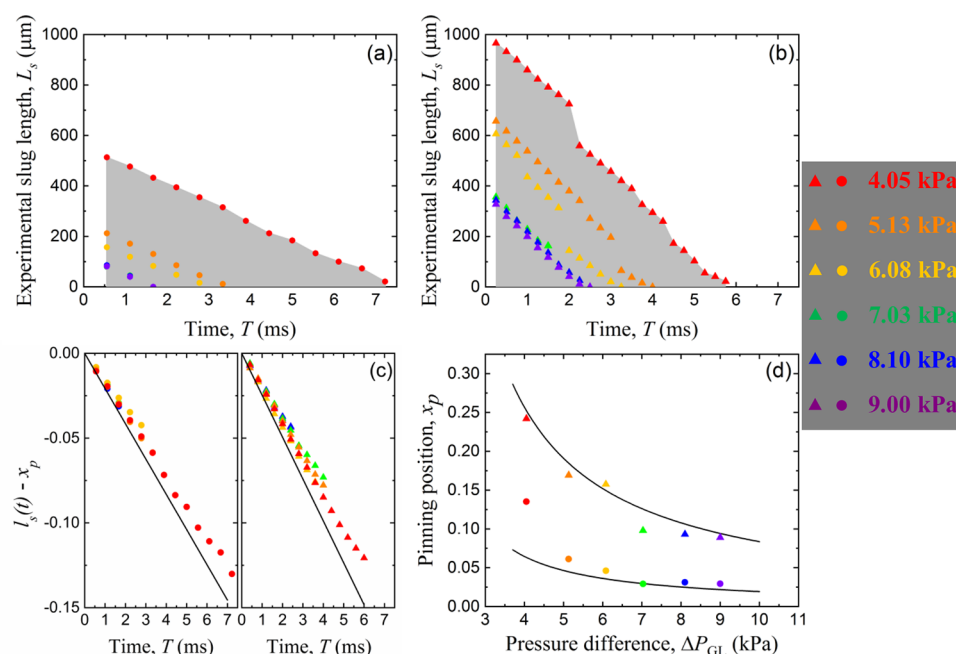


Figure 6. (a, b) Liquid slug lengths ($L_s = X_a - X_r$) during stable separation in the A-1 device at a liquid flow rate $Q_L = 20 \mu\text{L}/\text{min}$ (circle markers) and $90 \mu\text{L}/\text{min}$ (triangle markers) with a gas flow rate $Q_G = 350 \mu\text{L}_n/\text{min}$. The outlet pressure difference ΔP_{GL} was varied as follows: 4.05 kPa (red), 5.13 kPa (orange), 6.08 kPa (yellow), 7.03 kPa (green), 8.1 kPa (blue), and 9.0 kPa (violet). Although the rate of decrease of slug lengths remained constant, the pinning distance of the advancing meniscus decreased with increasing ΔP_{GL} . (c) Subtracting the experimentally measured slug lengths from the pinning position (markers) for $Q_L = 20 \mu\text{L}/\text{min}$ (left) and $90 \mu\text{L}/\text{min}$ (right) yields a linear behavior (solid lines), where $l_s(t)$ is the dimensionless length of the liquid slugs. (d) Pinning positions of the advancing meniscus (lines) compared with that of the experiments (markers) for two flow rates, $Q_L = 20 \mu\text{L}/\text{min}$ (circle markers) and $90 \mu\text{L}/\text{min}$ (triangle markers) at $Q_G = 350 \mu\text{L}_n/\text{min}$. The lines were obtained by fitting eq 14 to the experimental data, where β was used as the fitting parameter. For this device, $\beta = 2.5 \times 10^{-4}$ was obtained from the pinning position (x_p) vs ΔP_{GL} plots.

Slug Flow Characteristics: Stable Separation. High-speed imaging was performed to investigate gas–liquid separation in the A-1 device, as described earlier. At stable separation pressures (stage I), the front end of the liquid slug (advancing meniscus) was observed to be pinned inside the capillary zone (X_a is constant in Figure 5), where all the capillaries downstream were inactive for liquid separation. Such pinning behavior has been observed previously.^{17–19}

The image analysis algorithm was used to track the positions of the menisci from the point of entry of the receding menisci (i.e., from the third frame in Figure 3), and the liquid slug lengths and number of active capillaries were calculated (Figure 5 and Supplementary Movie 1). We observed that in the separation zone, the advancing meniscus is pinned, whereas the receding meniscus approached it linearly, as suggested by eq 15. It is also worth noting that, because of the curvature of the menisci, the number of active capillaries initially remained constant after the cap of the receding meniscus entered the separation zone (see dashed lines in Figure 5b). Similarly, when the two meniscus caps touched each other (hence, achieving complete separation, i.e., the eighth frame of Figure 3), both the distance between the two menisci at the edges of the capillaries and the number of active capillaries remained larger than zero due to the curvature of the menisci.

Varying the pressure difference ΔP_{GL} during separation showed that the rate of decrease in the lengths of the liquid slugs ($L_s = X_a - X_r$) remained constant (slopes of the datasets in Figures 6a,b), while only their position inside the separation zone varied, that is, the pinning positions (y intercepts in Figure 6a,b). Figure 6c depicts experimental data of the slug length normalized with the separation length (L) for different

values of ΔP_{GL} . It was observed that for a given flow rate, all data collapse into the same linear behavior as predicted by eq 15. It is worth noting that such linear behavior is independent of variation in ΔP_{GL} and only depends on the flow rate. Also, we note that the good agreement observed between the experiments and the theoretical behavior given by eq 15 indicates that the assumption made in the model, namely, the negligible slip velocity of the gas bubble inside the separation zone, is justified. The experiments in Figure 6d indicated that the slug became pinned further inside the capillary zone with decreasing ΔP_{GL} . For estimating the pressure drop along each capillary, ΔP_c in eq 8, which affects the pinning position (x_p), the capillary pressure of the advancing interface in the main channel (P_c) was required. P_c was estimated from the Young–Laplace equation of the form as in eq 1 and required the contact angle of the advancing meniscus and the surface tension of the liquid. Quantification of the apparent contact angle of the advancing menisci was a challenge due to the presence of droplet coalescence of the liquid slug with the droplets accumulated on the channel surfaces (due to their hydrophilicity) that led to changes in the curvature of the menisci. Various estimations exist in the literature obtained by either measuring the static contact angle in a pore³⁶ or considering the dependence of dynamic contact angles on the fluid velocity.^{37–39} Here, we took an equilibrium contact angle of 20° (air–water–glass combination³⁶) for the advancing meniscus to calculate P_c .

Equation 14 was fitted to the experimentally measured pinning positions of the advancing meniscus as shown in Figure 6d. For the data fit, β , which depends on the capillary geometry, was used as the fitting parameter. The value was

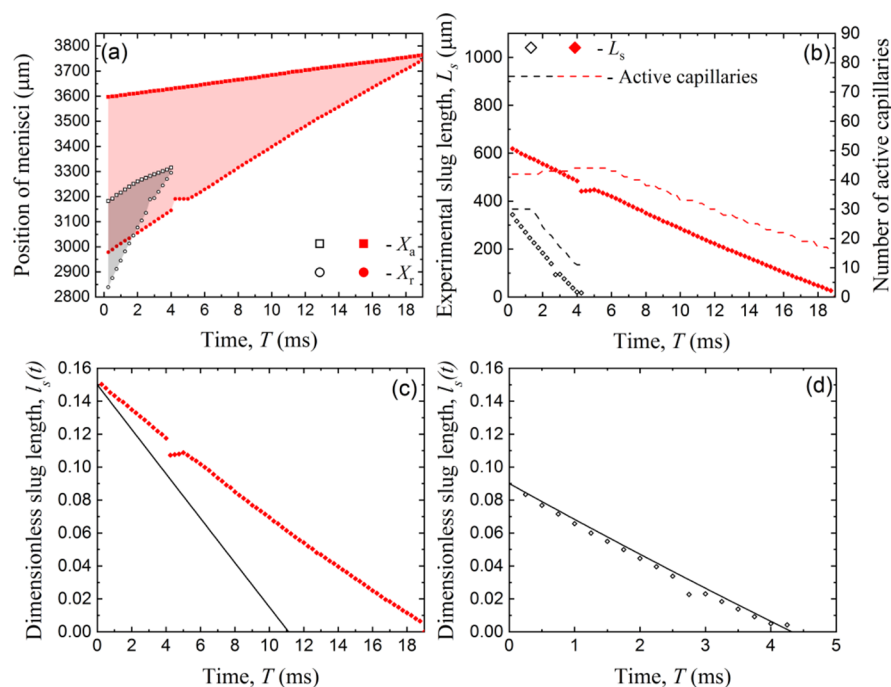


Figure 7. Tracking of liquid slugs at critical conditions in the A-1 device (i.e., last slug to completely separate before liquid breakthrough). (a) Positions of the advancing (square markers, X_a) and receding (circle markers, X_r) menisci from the first capillary. The graph shows the motion of the receding meniscus toward the nonstatic advancing meniscus as liquid separates through the capillaries. (b) The diamond markers show the length of the liquid slugs ($L_s = X_a - X_r$) within the separation zone measured between the meniscus caps. Dashed lines denote the number of active capillaries as the receding meniscus approaches the advancing meniscus. The liquid flow rate $Q_L = 20$ μL/min (red, closed markers) at an outlet pressure difference $\Delta P_{GL,critical} = 0.78$ kPa and 90 μL/min (black, open markers) at $\Delta P_{GL,critical} = 2.62$ kPa; both with a constant gas flow rate $Q_G = 350$ μL/min. (c, d) Comparison of the dimensionless slug length, $l_s(t)$, experimentally measured (markers) with eq 20 (solid lines) at (c) $Q_L = 20$ μL/min and (d) $Q_L = 90$ μL/min.

obtained by minimizing the error in the least-squares fit, considering both the data sets in Figure 6d simultaneously. For this particular A-1 device, we obtained $\beta = 2.5 \times 10^{-4}$. The behavior of the advancing meniscus before getting pinned is shown in Figure S6 of the Supporting Information. The advancing meniscus, $x_a(t)$, entered the separation zone and gradually reached x_p through an exponential behavior as described by eq 11. However, coalescence of water droplets in the main channel (formed due to the hydrophilic nature of the channel surfaces) with the advancing meniscus not only affected its curvature but also led to a variability in the pinning positions. This further exemplifies the difficulty in fully characterizing the liquid slug in the separation zone. The entire timescale of the separation process is shown in Figure S7 of the Supporting Information, highlighting the fluctuations in the positions of the advancing meniscus at high liquid flow rates through the sudden movement of the meniscus just before the receding meniscus approached it.

Once β was determined based on the capillary geometry, eqs 20 and 21 could be used to directly estimate the dynamics of the menisci of the slugs and the liquid-to-gas breakthrough pressures, as discussed below.

Slug Flow Characteristics: Critical Conditions. As the pressure difference ΔP_{GL} approached the liquid-to-gas breakthrough point, the pinning effect of the advancing meniscus was no longer observed, and the liquid slug moved further into the separation zone closer to the gas outlet. We define this as the critical point just before liquid-to-gas breakthrough, that is, when a liquid slug gets completely separated just before reaching the end of the separation zone. Owing to the sensitivity of the system, the slugs following the one at the

critical point were more likely to breakthrough into the gas outlet, and the system could not be maintained at the critical ΔP_{GL} . The flow of liquid through the capillaries took place also at the liquid-to-gas breakthrough pressure. Figure 7a,b shows the movement of the menisci, as the advancing meniscus approaches the gas outlet ($L = 4115$ μm for the A-1 device) and the (decreasing) lengths of the slugs at the critical condition. At this state, both menisci were moving at different speeds, leading to a nonlinear decrease in the liquid slug length. The liquid slug lengths were normalized with the separation length (L) and compared with the theoretical values given by eq 20 as shown in Figure 7c,d using the parameter β obtained from the previous section. We note that the observed discrepancy in Figure 7c could be a consequence of the aforementioned pressure sensitivity of the system.

During the liquid-to-gas breakthrough, the receding meniscus also reaches the gas outlet, and eq 21 applies. Figure 8 shows the experimental liquid-to-gas breakthrough pressures for different flow rates compared to the theoretical estimation given by eq 21. Castell et al.²³ hypothesized that a decrease in the advancing meniscus contact angle of the liquid slug with increasing flow rates and also when the meniscus entered the capillary zone was the potential reason for an increase in the breakthrough pressures with increasing flow rates. Similar behavior was also visually observed from the images of liquid slugs in the A-1 device.

Figure 8 also shows the experimental gas-to-liquid breakthrough compared to the Young–Laplace prediction, eq 1, hence providing the full operating zone of the separator. Experimental results of gas breakthrough show strong deviations from the Young–Laplace equation particularly at

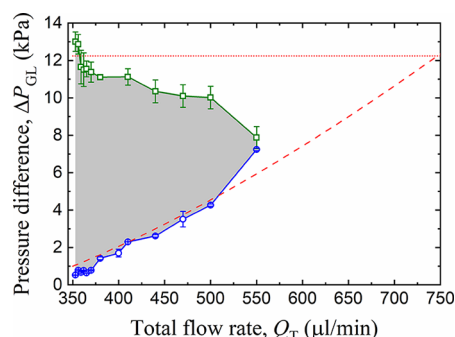


Figure 8. Operating zone of the gas-liquid separation device, A-1. Green square markers: points of gas-to-liquid breakthrough ($\Delta P_{G>L}$); blue circle markers: liquid-to-gas breakthrough ($\Delta P_{L>G}$). Dotted line: theoretical Young-Laplace breakthrough pressure through the capillaries; dashed line: liquid-to-gas breakthrough pressure given by eq 21. $\Delta P_{L>G}$ from the model was obtained from the dimensionalization of $\Delta p_{L>G}$ in the following form: $\Delta P_{L>G} = (P_0 \times \Delta p_{L>G}) + P_c$. The gas flow rate (Q_G) was 350 $\mu\text{L}/\text{min}$.

high flow rates, indicating that such theoretical approach may no longer be valid, thus necessitating a detailed investigation of the apparent contact angles of the menisci inside the capillaries.

Phase Separation in Separator A-2. To further investigate the ability of the model given by eq 21 to predict liquid-to-gas breakthrough, we performed additional gas-liquid separation studies in a different microseparator device, which we denote as A-2 (see Table S1 of the Supporting Information). The water-nitrogen-segmented flow was separated using this device via the same procedure adopted before. Characteristics of the slug flow in the separation zone were analyzed at two different flow rates and varying ΔP_{GL} (see Figure S8 of the Supporting Information). The events occurring during separation were found in general agreement with the model, namely, the linearity in the decrease of slug length and variation in the position of pinning of the advancing meniscus. The model estimated the pinning positions in the A-2 device for both the flow rates as shown in Figure 9a from eq 14 by fitting the parameter β , as described before, to a value of 2.1×10^{-4} . This value was then used to calculate the liquid-to-gas breakthrough pressures in the A-2 device as shown in

Figure 9b, which was compared with experimentally measured breakthrough values. The fitted β values were observed to be similar for the A-1 and A-2 devices, as the cross-sectional geometry of both the capillary designs was similar but with different number of capillaries and channel depths. In the A-2 device, the capillaries were farther apart ($R = 45 \mu\text{m}$, see Table S1 in the Supporting Information) than that of the A-1 device ($R = 25 \mu\text{m}$). Although the pressure drop across a single capillary is expected to be similar between the two devices, the fewer number of capillaries in the A-2 device ($N = 160$) resulted in a lower β value, thus leading to a lower outflow of liquid through the capillary array in comparison with that of the A-1 device with $N = 276$. For capillaries with nonparallel geometry, no empirical equations currently exist, and because of the tapered geometry, a larger flow rate for a given pressure drop is anticipated in comparison with a capillary with a parallel geometry of the same length and a width equal to the capillary width facing the main channel of the tapered geometry (D_i). This property makes β to be all the more sensitive.

We note that there is a good agreement up to values of $Q_T = 550 \mu\text{L}/\text{min}$ (Figure 9b) above which the theoretical estimations deviate from the experimental results. The deviations occur for $Q_L/Q_G > 0.5$ in the A-2 device, which has deeper channels. This led to shorter slug lengths, and as the liquid flow rate increased, the gas bubble lengths decreased. Under these conditions, a second liquid slug can enter the separation zone before the leading liquid slug has been fully separated through the capillaries. Because of this, the overall speed of the leading slug decreases hence giving rise to a lower liquid-to-gas breakthrough pressure differences. This effect is not captured by eq 21 and explains the deviation between theory and experimental results of Figure 9b for $Q_T > 550 \mu\text{L}/\text{min}$. Additionally, the deviations may also be attributed to the errors in the manual observations of the breakthrough pressures at such high flow rates along with the variations in the etch depths in the device resulting from the DRIE process.

CONCLUSIONS

We have presented a detailed experimental study on the separation process of a gas-liquid segmented flow in capillary microseparators that have an array of capillaries on both sides

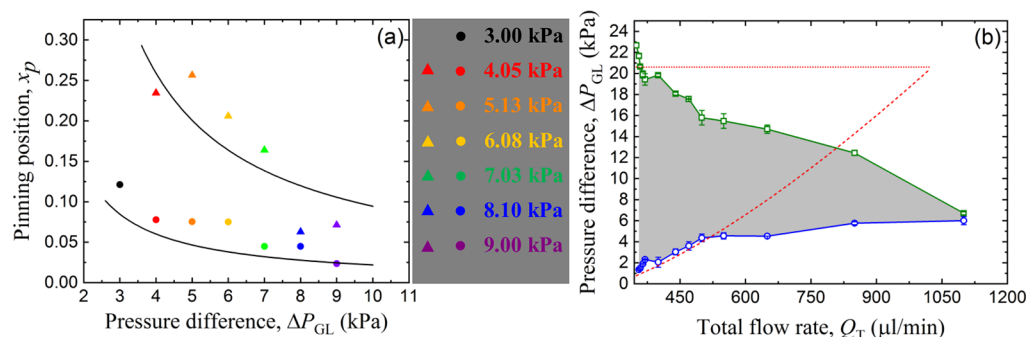


Figure 9. (a) Pinning positions of the advancing meniscus (lines obtained by fitting eq 14 with $\beta = 2.1 \times 10^{-4}$) compared with that of the experiments (markers) for two liquid flow rates, $Q_L = 20 \mu\text{L}/\text{min}$ (circle markers) and $90 \mu\text{L}/\text{min}$ (triangle markers), at a gas flow rate $Q_G = 350 \mu\text{L}/\text{min}$. The outlet pressure difference ΔP_{GL} was varied as follows: 3 kPa (black), 4.05 kPa (red), 5.13 kPa (orange), 6.08 kPa (yellow), 7.03 kPa (green), 8.1 kPa (blue), and 9.0 kPa (violet). (b) Liquid-to-gas breakthrough pressures ($\Delta P_{L>G}$) obtained from eq 21 (dashed line) and experimentally measured values (blue circle markers) for the A-2 device. Green square markers: experimental gas-to-liquid breakthrough pressures ($\Delta P_{G>L}$); horizontal dotted line: theoretical gas-to-liquid breakthrough pressures estimated from the Young-Laplace equation. $\Delta P_{L>G}$ from the model was obtained from the dimensionalization of $\Delta p_{L>G}$ in the following form: $\Delta P_{L>G} = (P_0 \times \Delta p_{L>G}) + P_c$.

of a channel. By making use of in situ optical high-speed imaging, we have been able to quantify the dynamics of the gas–liquid flow inside the separation zone. We observed that at stable separation conditions, the advancing meniscus became pinned in the separation zone, whereas the receding meniscus approached it at a constant speed, giving rise to a linear decrease of the liquid slug length. On the other hand, when the system approached critical conditions, both menisci were moving at different speeds, leading to an exponential decrease in the liquid slug length.

A hydrodynamic model based on mass conservation was developed, which estimated a linear decrease of the slug length under stable separation and an exponential decrease under critical conditions (just before liquid-to-gas breakthrough) in agreement with experiments. The model had only one fitting parameter, the geometric factor β , which was obtained from the location of the pinned position of the advancing meniscus as a function of the pressure difference between the gas outlet and the liquid outlet. Once this parameter was determined, the model predicted well the liquid-to-gas breakthrough pressures. The model was further tested against the performance of another microseparation device with different geometrical design of the capillaries and with different etch depth. The model estimated the liquid-to-gas breakthrough pressures with a good agreement up to $Q_T = 550 \mu\text{L}/\text{min}$, above which more than one slug was encountered in the separation zone, which was not captured in our model.

■ ASSOCIATED CONTENT

■ Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: [10.1021/acs.langmuir.8b04202](https://doi.org/10.1021/acs.langmuir.8b04202).

Design and operation of the microfabricated gas–liquid separation devices; image preprocessing via morphological reconstruction; measurement of the initial liquid slug length, number of active capillaries, and bubble frequency during separation; tracking the entry of the advancing meniscus in the separation zone at stable separation conditions; quantification of slug lengths in the A-2 device at varying pressure differences (PDF)

A movie of the image analysis algorithm tracking the gas–liquid menisci, recorded at 4000 fps and played back at 3 fps (AVI)

A video of gas–liquid separation occurring at $Q_L = 90 \mu\text{L}/\text{min}$, recorded at 4000 fps and every 50th frame played back at 10 fps (AVI)

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Author Contributions

The manuscript was written through contributions from all authors. All authors have given approval to the final version of the manuscript and declare no conflicting interests.

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

We are grateful to the anonymous reviewer for valuable comments and suggestions and to Dr. Benjamin Aymard and Mr. Urbain Vaes, Imperial College London, for valuable inputs and stimulating discussion. We acknowledge financial support from the Engineering and Physical Sciences Research Council of the U.K. through grant nos. EP/L027232 and EP/L027186.

■ NOTATION

Δp	dimensionless pressure drop, $(\Delta P_{GL} - \Delta P_c)/P_0$
$\Delta P_{G>L}$	ΔP_{GL} at which gas-to-liquid breakthrough occurs (kPa)
$\Delta p_{L>G}$	dimensionless liquid-to-gas breakthrough pressure obtained from eq 21,
$\Delta P_{L>G}$	ΔP_{GL} at which liquid-to-gas breakthrough occurs (kPa)
ΔP_c	pressure drop across a capillary (kPa)
ΔP_{GL}	gas–liquid outlet pressure difference, $P_G - P_L$ (kPa)
A	cross-sectional area of the channel (μm^2)
A_c	cross-sectional area of a capillary (μm^2)
D	capillary width (μm)
H	channel/capillary height (μm)
L	length of the separation zone (mm)
l_0	dimensionless initial length of the liquid slug, L_0/L
L_0	initial length of the liquid slug (mm)
L_c	length of capillaries (μm)
l_s	dimensionless length of the liquid slug, $x_a - x_r$
L_s	length of the liquid slug in the separation zone, $X_a - X_r$ (μm)
M	mass of the part of the slug in contact with the microcapillaries (kg)
M_{outflow}	mass outflow of liquid through the capillaries (kg)
N	total number of capillaries
P_0	pressure scale for nondimensionalization, $\eta U_0/L_c$
P_a	liquid-side pressure at the advancing interface (kPa)
P_c	capillary pressure of the advancing interface (kPa)
P_G	gas outlet pressure (kPa)
P_{in}	total inlet pressure (kPa)
P_L	liquid outlet pressure (kPa)
P_r	liquid-side pressure at the receding interface (kPa)
Q_c	local flow rate through each capillary ($\mu\text{L}/\text{min}$)
Q_G	gas flow rate ($\mu\text{L}/\text{min}$)
q_L	dimensionless liquid flow rate, Q_L/Q_G
Q_L	liquid flow rate ($\mu\text{L}/\text{min}$)
q_T	dimensionless total flow rate, Q_T/Q_G
Q_T	total flow rate ($\mu\text{L}/\text{min}$)
t	dimensionless time scale, T/T_0
T	time (ms)
T_0	time scale for non-dimensionalization, L/U_0
U	superficial average velocity of the two-phase flow (mm/s)
U_0	velocity term for non-dimensionalization, Q_G/A
W	width of main channel (μm)
x_a	dimensionless advancing meniscus position, X_a/L
X_a	position of the advancing meniscus (μm)
x_p	pinning position of the advancing meniscus
x_r	dimensionless receding meniscus position, X_r/L
X_r	position of the receding meniscus (μm)

■ GREEK LETTERS

α geometrical correction factor in the Hagen–Poiseuille equation

- β modified geometrical factor presented in this work = $NA_c^2 / \alpha AL_c^2$
 γ surface tension of the liquid (N/m)
 η viscosity of the liquid (Pa·s)
 θ_e equilibrium contact angle ($^\circ$)
 ρ density of the liquid (kg/m³)

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